AUG. 10. 2006 11:19AM HITT GAINES 9724808865 NO. 5922 P. 1/16

HITT GAINES, P.C.

Intellectual Property Law & Related Matters

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USPTO

Examiner Chukwuma O. Nwaonicha; Art Unit 1621

FAX NO.

(571) 273-8300

AUG 1-1 2006 of Correction

FROM:

Ronald J. Corbett

RE:

Patent No. 7,012,140 issued March 14, 2006

Attorney Docket No.: UTAD-0004

DATE:

August 10, 2006

PAGES:

16 (including cover page)

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Documents transmitted herewith:

facsimile transmittal (pp.1);

Certificate of Correction Transmittal Letter (pp.6); and

Certificate of Correction (pp.9).

DOCKET NO. UTAD-0004

RECEIVED CENTRAL FAX CENTER

PATENT

AUG 1 0 2006

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re U.S. Registered Patent No. 7,012,140 of:

A. Dean Sherry, Mark Woods and Zoltan Kovacs

Issued:

March 14, 2006

Filed: July 14, 2003

Serial No.:

10/619,362

For:

SELECTION OF COORDINATION GEOMETRY TO ADJUST WATER

EXCHANGE RATES OF PARAMAGNETIC METAL ION-BASED

MACROCYCLIC CONTRAST AGENTS

Group:

1621

Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

> I hereby certify that this correspondence is being facsimils transmitted to the Patent and Trademark Office (Fax No. (571) 273-8300) on August 10, 2006

erson singing the certificate

Signature of person mailing

Sir:

CERTIFICATE OF CORRECTION TRANSMITTAL LETTER

The following clerical or typographical errors were noted in Column 4, lines 14-17:

FIGURE 3A and FIGURE 3B illustrates exemplary ¹H NMR spectra of Eu(S RRRR- $NO_2BnDOMA)$ [Eu(S-RRRR-NO_2BnDOTMA)] and [Eu(S-SSSS-NO_2BnDOTMA)], respectively, produced according to the present; and

The following clerical or typographical errors were noted in Column 5, lines 60-67 through and including Column 6, lines 1-3:

For example, when the three or more pendant arm carbon atoms C' have Δ orientations and the chirality of the one or more R^6 –substituted ring carbons is selected such that the macrocyclic ring has an identical (8888) orientation, then the tetraazacyclododecane ligand has a capped twisted square antiprism configuration. Or, when the three or more pendant arm carbon atoms C' have Λ orientations arid and the chirality of the one or more ring carbons is selected such that the macrocyclic ring has a (λλλλ) orientation, then the tetraazacyclododecane ligand again has a capped twisted square antiprism configuration.

The following clerical or typographical errors were noted in Column 8, lines 61-67 through and including Column 9, lines 1-11:

In certain preferred embodiments of the method 100, the contrast agent further includes a carrier component, conjugated to one or more of the functionalized substituents R6, as discussed above. In certain embodiments of the present invention, the CA includes at least one and up to twenty of the tetraazacyclododecane ligands. Such ligands may be covalently or noncovalently bonded to a carrier component, such as described above, comprising a portion of the contrast agent. Collecting several such ligands, and associated metal ions and bound water molecules, allows a larger effective magnetic resonance signal to be achieved at lower concentrations of contrast agent. In certain such embodiments, where the water molecule (H2O) associated with the tetraazacyclododecane ligand has a $\tau_{\rm M}^{298}$, of between about 10 and about 100 nanoseconds, the water molecule associated with a contrast agent that further includes a carrier component has a relativity relaxivity at 298°C, r₁²⁹⁸, of at least about 50 mM⁻¹ s⁻¹.

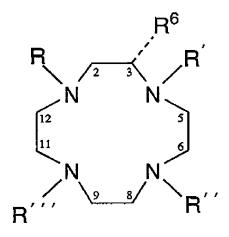
The following clerical or typographical errors were noted in Column 11, lines 59-67 through and including Column 12, lines 1-5:

The two complexes $[Gd(S-RRRR-NO_2BnDOTMA)]^-$ and $[Gd(s-ssss-N)_2BnDOTMA)]^ [Gd(S-SSSS-NO_2BnDOTMA)]^-$ have substantially different $1/T_2$ - temperature profiles, indicating substantially different water exchange rates. The profile for $[Gd(S-RRRR-N)_2BnDOTMA)]^ [Gd(S-RRRR-NO_2BnDOTMA)]^-$, rises, maximizes and then falls away with increasing temperature, indicative of fairly slow water exchange. In contrast, the profile for $[Gd(S-SSSS-NO_2BnDOTMA)]^-$, did not reach a maximum within the temperature range study, indicative of a more rapid water exchange. The values of τ_M^{298} obtained by fitting procedures, well known to those skilled in the art, to profiles such as depicted in FIGURE 4, are summarized in TABLE 2. Also shown in TABLE 2 are the relaxivities of water molecules associated with these semens isomers at 25 and 37 °C.

The following clerical or typographical error was noted in Claim 1:

1. A magnetic resonance contrast agent compound comprising:

a tetraacyclododecane tetraazacyclododecane ligand having a general structural formula as follows:



and comprising a macrocyclic ring and wherein pendant arms R, R', R' and R'' attached to a ring nitrogen have the general formula: -C'HR'R² and for three or more of said pendant arms a chirality of said carbon atoms C' are identical for each of said three or more pendant arms, said R¹ are groups larger than hydrogen, and said R² is selected from the goup group consisting of:

```
an alcohol (-CH<sub>2</sub>OH);
amides (-CONR<sup>3</sup>R<sup>4</sup>, where R<sup>3</sup> and R<sup>4</sup> are organic groups);
a carboxylate (-COOH);
phosphinates (-PO<sub>2</sub>HR<sup>5</sup>, where R<sup>5</sup> is an organic group); and
a phosphonate (-PO(OH)2); and
```

wherein one or more of substituents R⁶ is a group larger than a methyl group and is located on one or more ring carbons; and

a paramagetie paramagnetic metal ion coordinated to said tetraazacyclododecane ligand.

The following clerical or typographical error was noted in Claim 5:

5. The magnetic resonance contrast agent compound as recited in Claim 1, wherein said chirality of said carbon atoms C' is controlled to provide said three or more of said pendant arms with a Δ or Λ orientation, and wherein a chirality of a ring carbon bonded to said one or more of substituents R⁶ provides said macrocyclic macrocyclic ring with an opposite orientation, xxxx or δδδδ, respectively, said tetraazacyclododecane ligand thereby having a monocapped square antiprism coordination geometry.

The following clerical or typographical error was noted in Claim 6:

6. The magnetic resonance contrast agent compound as recited in Claim 5, wherein said R² group is said alcohol or amide, and further including a water molecule associated with said tetraazacyclododecane ligand and said paramagnetic paramagnetic metal ion, said water molecule having a residence lifetime at about 298°K, $\tau_{\rm M}^{298}$, of between about 10 and about 5000 microseconds.

The following clerical or typographical error was noted in Claim 7:

7. The magnetic resonance contrast agent compound as recited in Claim 5, wherein said R² group is said carboxyl, and further including a water molecule associated with said tetreyclododecane tetraazacyclododecane ligand and said paramagnetic metal ion, said water molecule having a residence lifetime at about 298°K, $\tau_{\rm M}^{298}$, of between about 100 and about 500 nanoseconds.

The following clerical or typographical error was noted in Claim 8:

8. The magnetic resonance contrast agent compound as recited in Claim 5, wherein said R² group is said phosphonate or said phosphinate, and further including a water molecule associated with said tet azacyclododecane tetraazacyclododecane ligand and said paramagnetic metal ion, said water molecule having a residence lifetime at about 298°K, $\tau_{\rm M}^{298}$, of between about 10 and about 100 nanoseconds.

The following clerical or typographical error was noted in Claim 9:

9. The magnetic resonance contrast agent compound as recited in Claim 1, wherein said R¹ is a methyl group, said R² is said earboxyl carboxyl, and said R⁶ is a para-aminobenzyl group and said paramagnetic metal ion is Gd³⁺.

The following clerical or typographical error was noted in Claim 11:

11. The magnetic resonance contrast agent compound as recited in Claim 1, wherein at least one of said one or more of substituents R⁶ include a functional group selected from the group consisting of:

amino groups;

carboxylates;

isothiocyanates; and

maleiimdes; and

a carrier component conjugated to said functional group.

Applicants respectfully request that a Certificate of Correction be issued pursuant to 37 C.F.R. §1.322. A Certificate of Correction reflecting these changes is enclosed. There is no fee associated with this correction since the errors listed above were made by the USPTO. However, the Commissioner is authorized to credit any overages or charge any additional fees in connection with this communication to Deposit Account Number 08-2395.

Respectfully submitted,

HITT GAINES, P.C.

Ronald J. Corbett

Registration No. 47,500

Date: August 10, 2006

Hitt Gaines, P.C. P.O. Box 832570 Richardson, Texas 75083-2570 (972) 480-8800

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UNITED STATES PATENT AND TRADEMARK OFFICE CERTIFICATE OF CORRECTION

PATENT NO. : 7,012,140

DATED

: March 14, 2006

INVENTOR(S): A. Dean Sherry, Mark Woods and Zoltan Kovacs

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

Column 4, lines 14-17 should appear as follows:

FIGURE 3A and FIGURE 3B illustrates exemplary 1H NMR spectra of [Eu(S-RRRR-NO2BnDOMA)] [Eu(S-RRRR-NO2BnDOTMA)] and [Eu(S-SSSS-NO₂BnDOTMA)], respectively, produced according to the present; and

Column 5, lines 60-67 through and including Column 6, lines 1-3 should appear as follows:

For example, when the three or more pendant arm carbon atoms C' have Δ orientations and the chirality of the one or more R⁶-substituted ring carbons is selected such that the macrocyclic ring has an identical ($\delta\delta\delta\delta$) orientation, then the tetraazacyclododecane ligand has a capped twisted square antiprism configuration. Or, when the three or more pendant arm carbon atoms C' have Λ orientations axid and the chirality of the one or more ring carbons is selected such that the macrocyclic ring has a (λλλλ) orientation, then the tetraazacyclododecane ligand again has a capped twisted square antiprism configuration.

MAILING ADDRESS OF SENDER:

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7,012,140

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UNITED STATES PATENT AND TRADEMARK OFFICE CERTIFICATE OF CORRECTION

PATENT NO.

: 7.012.140

DATED

: March 14, 2006

INVENTOR(S): A. Dean Sherry, Mark Woods and Zoltan Kovacs

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

Column 8, lines 61-67 through Column 9, lines 1-11 should appear as follows:

In certain preferred embodiments of the method 100, the contrast agent further includes a carrier component, conjugated to one or more of the functionalized substituents R⁶, as discussed above. In certain embodiments of the present invention, the CA includes at least one and up to twenty of the tetraazacyclododecane ligands. Such ligands may be covalently or noncovalently bonded to a carrier component, such as described above, comprising a portion of the contrast agent. Collecting several such ligands, and associated metal ions and bound water molecules, allows a larger effective magnetic resonance signal to be achieved at lower concentrations of contrast agent. In certain such embodiments, where the water molecule (H2O) associated with the tetraazacyclododecane ligand has a TM²⁹⁸, of between about 10 and about 100 nanoseconds, the water molecule associated with a contrast agent that further includes a carrier component has a relativity relaxivity at 298°C, r1²⁹⁸, of at least about 50 mM⁻¹ s⁻¹.

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7,012,140

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UNITED STATES PATENT AND TRADEMARK OFFICE CERTIFICATE OF CORRECTION

PATENT NO. : 7,012,140

DATED

: March 14, 2006

INVENTOR(S): A. Dean Sherry, Mark Woods and Zoltan Kovacs

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

Column 11, lines 59-67 through Column 12, lines 1-5 should appear as follows:

The two complexes [Gd(S-RRRR- NO2BnDOTMA)] and [Gd(s-ssss- N)2BnDOTMA)] [Gd(S-SSSS-NO2BnDOTMA)] have substantially different 1/T₂ temperature profiles, indicating substantially different water exchange rates. The profile for [Gd(S-RRRR-N)2BnDOTMA)] [Gd(S-RRRR-NO2BnDOTMA)], rises, maximizes and then falls away with increasing temperature, indicative of fairly slow water exchange. In contrast, the profile for [Gd(S-SSSS-NO₂BnDOTMA)], did not reach a maximum within the temperature range study, indicative of a more rapid water exchange. The values of τM^{298} obtained by fitting procedures, well known to those skilled in the art, to profiles such as depicted in FIGURE 4, are summarized in TABLE 2. Also shown in TABLE 2 are the relaxivities of water molecules associated with these somers at 25 and 37 °C.

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UNITED STATES PATENT AND TRADEMARK OFFICE CERTIFICATE OF CORRECTION

PATENT NO.

7,012,140

DATED

: March 14, 2006

INVENTOR(S) : A. Dean Sherry, Mark Woods and Zoltan Kovacs

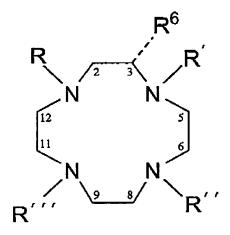
It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

Claim 1 should appear as follows:

The following clerical or typographical error was noted in Claim 1:

1. A magnetic resonance contrast agent compound comprising:

a tetraacyclododecane tetraazacyclododecane ligand having a general structural formula as follows:



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UNITED STATES PATENT AND TRADEMARK OFFICE CERTIFICATE OF CORRECTION

PATENT NO. : 7,012,140

DATED

: March 14, 2006

INVENTOR(S): A. Dean Sherry, Mark Woods and Zoltan Kovacs

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

(claim 1 continued)

and comprising a macrocyclic macrocyclic ring and wherein pendant arms R, R', attached to a ring nitrogen have the general formula: -C'HR'R2 and for three or more of said pendant arms a chirality of said carbon atoms C' are identical for each of said three or more pendant arms, said R¹ are groups larger than hydrogen, and said R² is selected from the goup group consisting of:

an alcohol (-CH2OH);

amides (-CONR³R⁴, where R³ and R⁴ are organic groups);

a carboxylate (-COOH);

phosphinates (-PO2HR⁵, where R⁵ is an organic group); and

a phosphonate (-PO(OH)2); and

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PATENT NO. : 7,012,140

DATED

: March 14, 2006

INVENTOR(S): A. Dean Sherry, Mark Woods and Zoltan Kovacs

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

(claim 1 concluded)

wherein one or more of substituents R⁶ is a group larger than a methyl group and is located on one or more ring carbons; and

a paramagetie paramagnetic metal ion coordinated to said tetraazacyclododecane ligand.

The correction to claim 1 is fully supported in Column 12, lines 48, 61 and 67 and Column 13, line 11.

Claim 5 should appear as follows:

5. The magnetic resonance contrast agent compound as recited in Claim 1, wherein said chirality of said carbon atoms C' is controlled to provide said three or more of said pendant arms with a Δ or Λ orientation, and wherein a chirality of a ring carbon bonded to said one or more of substituents R⁶ provides said macrocyclic macrocyclic ring with an opposite orientation, λλλλ or δδδδ, respectively, said tetraazacyclododecane ligand thereby having a monocapped square antiprism coordination geometry.

The correction to claim 5 is fully supported in Column 13, line 41.

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7,012,140

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PATENT NO. : 7,012,140

DATED

: March 14, 2006

INVENTOR(S): A. Dean Sherry, Mark Woods and Zoltan Kovacs

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

Claim 6 should appear as follows:

6. The magnetic resonance contrast agent compound as recited in Claim 5, wherein said R² group is said alcohol or amide, and further including a water molecule associated with said tetrazzacyclododecane ligand and said paramagnetic paramagnetic metal ion, said water molecule having a residence lifetime at about 298°K, $\tau_{\rm M}^{298}$, of between between about 10 and about 5000 microseconds.

The correction to claim 6 is fully supported in Column 14, lines 2, 4, 5 and 6.

Claim 7 should appear as follows:

7. The magnetic resonance contrast agent compound as recited in Claim 5, wherein said R² group is said carboxyl, and further including a water molecule associated with said tetrayelododecane tetraazacyclododecane ligand and said paramagnetic metal ion, said water molecule having a residence lifetime at about 298°K, τ_M^{298} , of between about 100 and about 500 nanoseconds.

The correction to claim 7 is fully supported in Column 14, line 11.

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It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

Claim 8 should appear as follows:

8. The magnetic resonance contrast agent compound as recited in Claim 5, wherein said R² group is said phosphonate or said phosphinate, and further including a water molecule associated with said tet azacyclododecane tetraazacyclododecane ligand and said paramagnetic metal ion, said water molecule having a residence lifetime at about 298°K, τ_M^{298} , of between about 10 and about 100 nanoseconds.

The correction to claim 8 is fully supported in Column 14, lines 18 and 20.

Claim 9 should appear as follows:

9. The magnetic resonance contrast agent compound as recited in Claim 1, wherein said R^1 is a methyl group, said R^2 is said carboxyl carboxyl, and said R^6 is a para-aminobenzyl group and said paramagnetic metal ion is Gd³⁺.

The correction to claim 9 is fully supported in Column 14, line 24.

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7,012,140

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UNITED STATES PATENT AND TRADEMARK OFFICE CERTIFICATE OF CORRECTION

PATENT NO. : 7,012,140

DATED

: March 14, 2006

INVENTOR(S): A. Dean Sherry, Mark Woods and Zoltan Kovacs

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

Claim 11 should appear as follows:

11. The magnetic resonance contrast agent computed compound as recited in Claim 1, wherein at least one of said one or more of substituents R⁶ include a functional group selected from the group consisting of:

amino groups;

carboxylates;

isothiocyanates; and

maleiimdes; and

a carrier component conjugated to said functional group.

The correction to claim 11 is fully supported in Column 14, line 31.

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